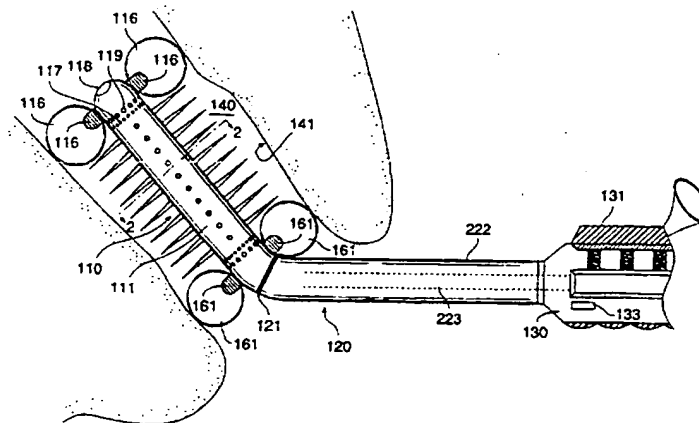




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(54) Title: ABLATION OF RECTAL AND OTHER INTERNAL BODY STRUCTURES

(57) Abstract

The invention provides an apparatus and system for ablation of body structures or tissue in the region of the rectum. A catheter is inserted into the rectum, and an electrode is disposed thereon for emitting energy. The environment for an ablation region is isolated or otherwise controlled by blocking gas or fluid using a pair of inflatable balloons at upstream and downstream locations. Inflatable balloons also serve to anchor the catheter in place. A plurality of electrodes are disposed on the catheter and at least one such electrode is selected and advanced out of the catheter to penetrate and ablate selected tissue inside the body in the region of the rectum. The electrodes are coupled to sensors to determine control parameters of the body structure or tissue, and which are used by feedback technique to control delivery of energy for ablation or fluids for cooling or hydration. The catheter includes an optical path disposed for coupling to an external view piece, so as to allow medical personnel to view or control positioning of the catheter and operation of the electrodes. The catheter is disposed to deliver flowable substances for aiding in ablation, or for aiding in repair of tissue, such as collagen or another substance for covering lesions or for filling fissures. The flowable substances are delivered using at least one lumen in the catheter, either from at least one hole in the catheter, from an area of the catheter covered by a microporous membrane, or from microporous balloons.

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SPECIFICATION**Title of the Invention****Ablation of Rectal and Other Internal Body Structures****Cross-Reference to Related Applications**

This application is a continuation-in-part of the following co-pending applications:

- 5 o Application Serial No. 08/677,811, entitled "Treating Aneurysms by Applying Hardening/Softening Agents to Hardenable/Softenable Substances," filed July 10, 1996, attorney docket number MOED-001;
- 10 o Application Serial No. 08651,800, entitled "Method and Apparatus for Treatment of Airway Obstruction," filed May 22, 1996, attorney docket number SOMN-1009-4;
- 15 o Application Serial No. 08/651,378, entitled "Method and Apparatus for Treatment of Airway Obstruction," filed May 22, 1996, attorney docket number SOMN-1009-5;
- 20 o Application Serial No. 08/643,203, entitled "Uvula, Tonsil, Adenoid and Sinus Tissue Treatment Device and Method," filed May 6, 1996, attorney docket number SOMN-1011-1;
- 25 o Application Serial No. 08/643,524, entitled "Ablation Apparatus and System for Removal of Soft Palate," filed May 6, 1996, attorney docket number SOMN-1012-2;
- 30 o Application Serial No. 08/651,796, entitled "Apparatus and Methods for Ablating Turbinates," filed May 22, 1996, attorney docket number SOMN-2001;
- o Application Serial No. 08/651,798, entitled "Apparatus and Methods for Ablating Turbinates," filed May 22, 1996, attorney docket number SOMN-2002;
- o Application Serial No. 08/660,539, entitled "Precision Placement of Ablation Apparatus," filed June 7, 1996, attorney docket number SOMN-2003; and

o Application Serial No. 08/663,004, entitled "Electrode for Ablating Tissue," filed June 7, 1996, attorney docket number SOMN-2004.

5 Each of these applications is hereby incorporated by reference as if fully set forth herein.

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Background of the Invention

1. Field of the Invention

This invention relates to ablation of rectal and other internal body structures.

15

2. Description of Related Art

Human beings are subject to a number of disorders in the area of the rectum and colon, including hemorrhoids (external and internal), prolapse of the rectal muscles, rectal muscle spasms, anal fissures, polyps, diverticulosis and diverticulitis, and pilonidal cysts. Other internal disorders in nearby regions of the body include (in men) prostate cancer, (in women) incontinence, vaginal bleeding, vaginal cysts, vaginal fibroids, prolapse of the uterus, and related tumors or cancerous tissue.

20

Although there are treatments available for these disorders, such as surgery, systemic or topical medication, these treatments suffer from various drawbacks, including (for surgery) their relative invasiveness and expense, and (for medicinal approaches) their relative ineffectiveness and the causation of serious side-effects. Accordingly, it would be advantageous to provide methods and apparatus for treatment which are not subject to the drawbacks of surgery and medicinal approaches.

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Although it is known to use RF energy to ablate tissue in the body (such as heart muscle tissue) to treat disorders, one problem which has arisen in the art is accounting for the flow of bodily fluids and gases while ablating tissue.

Bodily fluids can dissipate, and can detrimentally absorb, energy to be applied to tissue.

5 Accordingly, it would be advantageous to provide improved techniques for treatment of disorders in the area of the rectum and colon. This advantage is achieved by a method and system according to the present invention in which a catheter is inserted into the rectum, and at least one electrode is disposed thereon for emitting energy to ablate body structures or other tissue in an ablation region in or near the rectum, such as the sphincter, rectum, colon, or
10 prostate.

Summary of the Invention

15 The invention provides a method and system for ablation of body structures or tissue in an ablation region in or near the rectum (such as the sphincter, rectum, colon, or prostate). A catheter is inserted into the rectum, and at least one electrode is disposed thereon for emitting energy to ablate body structures or other tissue, such as by cell death, dehydration, or denaturation. The environment for the ablation region is isolated or otherwise controlled, such as by
20 blocking gas or fluid using a pair of inflatable balloons at upstream and downstream locations from the ablation region. In a preferred embodiment, inflatable balloons also serve to anchor the catheter in place and prevent the catheter from being expelled from the body.

25 In preferred embodiments, the catheter is flexible for reaching a selected internal organ or region, a plurality of electrodes are disposed on the catheter and at least one such electrode is selected and advanced out of the catheter to penetrate and ablate selected tissue inside the body in ablation region in or near the rectum, such as an individual cyst, hemorrhoid, polyp, tumor, or other selected
30 lesion or tissue. The electrodes are coupled to sensors to determine control parameters of the body structure or tissue, such as impedance or temperature, and which are used by feedback technique to control delivery of energy for ablation or fluids for cooling or hydration. In a preferred embodiment, the catheter includes an optical path disposed for coupling to an external view piece, so as to allow
35 medical personnel to view or control positioning of the catheter and operation of the electrodes.

5 In further preferred embodiments, the catheter is disposed to deliver flowable substances for aiding in ablation, such as saline or antibiotics, or for aiding in repair of tissue (either before or after ablation), such as collagen or another substance for covering lesions or for filling fissures in or near the ablation region, or for other medicinal effects, such as anesthetic, anti-inflammatory, or antispasmodic substances. The flowable substances are delivered using at least one lumen in the catheter, either from at least one hole in the catheter, from an area of the catheter covered by a microporous membrane, or from microporous balloons (either the same as or in addition to balloons used to anchor the catheter in place or to block gas or fluid).

Brief Description of the Drawings

15 Figure 1 shows a side view of a catheter and electrode assembly.

Figure 2 shows a cut-away view of a catheter, taken along a line 2--2 in Figure 1.

20 Figure 3 shows a method of treatment of a hemorrhoid.

Figure 4 shows a method of treatment of a prolapsed or spasmodic muscle.

25 Figure 5 shows a method of treatment of an anal fissure.

Figure 6 shows a method of treatment of a tumor in the prostate.

Description of the Preferred Embodiment

30 Catheter and Electrode Assembly

Figure 1 shows a side view of a catheter and electrode assembly.

35 An assembly 100 for ablating rectal and other internal body structures includes a catheter 110, a control and delivery linkage 120, and a control element 130.

The catheter 110 is coupled to the control and delivery linkage 120 using a gearing element 121, which allows the catheter 110 to be rotated with respect to the control and delivery linkage 120 by an operator using the control element 130.

5 The catheter 110 includes a base 111, having a substantially cylindrical shape, coupled at a proximal end to the gearing element 121, and having a distal end. The catheter 110 is preferably disposed for insertion into the rectum at an angle to the control and delivery linkage 120, preferably an angle between about 30° and about 45° less than a right angle. The catheter 110 is between about 1 inch (2.54
10 cm) and about 2 inches (5.08 cm) in diameter, and between about 6 inches (15.24 cm) and about 8 inches (20.32 cm) in length.

15 The catheter 110 includes a plurality of holes 112, and a plurality of electrodes 113 which may be extended from at least some of the holes 112. The holes 112 are spaced regularly around the circumference and along the length of the catheter 110, having a spacing of about 0.25 inches (0.64 cm) between adjacent holes 112. The electrodes 113 are spaced regularly to occupy about one-half of the holes 112, and are between about 0.5 cm and about 1.0 cm in length.

20 The electrodes 113 each include a metallic tube 114 defining a hollow lumen 115, shaped similarly to an injection needle, so as to be disposed to deliver at least one flowable substance to a region 140 near the catheter 110. In a preferred embodiment, the deliverable flowable substance includes saline with a concentration of less than about 10% NaCl, which aids in both hydration of body
25 structures and other tissue, and in delivery of RF energy to the region 140. However, in alternative embodiments, the deliverable flowable substance includes other substances, including saline with other concentrations, systemic or topical antibiotics, collagen or another hardenable substance, or other bioactive, chemoactive, or radioactive substances (including anesthetic, anti-inflammatory, or
30 antispasmodic substances, or tracer materials).

35 The catheter 110 includes at least one balloon 116, disposed for inflation so as to block gas or fluid from the body from entering the region 140. In a preferred embodiment, there is a distal balloon 116 disposed at the distal end of the catheter 110 and there is a proximal balloon 116 disposed at the proximal end of the catheter 110. The distal balloon 116 and the proximal balloon 116 preferably each

comprise ring-shaped balloons, disposed so that when inflated each surrounds the catheter 110 and makes a gas-tight or fluid-tight seal, both with the catheter 110 and with a wall 141 of the rectum or other body structure into which the catheter 110 is inserted. However, in alternative embodiments, the distal balloon 116 may
5 comprise a spherical or ellipsoidal balloon disposed at the distal end of the catheter 110 in such manner that when inflated it surrounds the catheter 110 and makes a gas-tight or fluid-tight seal with the wall 141.

The catheter 110 also includes at least one balloon 116 disposed to anchor the
10 catheter 110 at a selected location within the rectum or other body structure into which the catheter 110 is inserted. In a preferred embodiment, the balloon 116 used to anchor the catheter 110 is the proximal balloon 116, which when inflated prevents the catheter 110 from being expelled from the body in like manner as the operation of a Foley catheter. However, in alternative embodiments, the balloon
15 116 used to anchor the catheter 110 may comprise an additional or alternative balloon which is disposed solely or primarily for the purpose of anchoring the catheter 110 into its selected place, again in like manner as the operation of a Foley catheter.

20 The catheter 110 includes a fluid circulation system 117, including at least one fluid outlet port and at least one fluid inlet port. The fluid circulation system 117 is disposed for providing fluid in the region near the catheter 110, such as for delivering fluid for cooling the region 140 and for removing other fluid for aspirating the region 140.

25 The catheter 110 includes an optical view port 118, possibly including a lens or other transparent or translucent covering, disposed to allow inflow of light (visible or infrared) for transmission to an operator for viewing and control of the operation of the catheter 110.

30 The catheter 110 includes at least one sensor 119, such as a sensor 119 for impedance or temperature. In a preferred embodiment, the temperature sensor 119 includes a thermocouple, but in alternative embodiments, the temperature sensor 119 may include a thermistor or other device for sensing temperature and
35 providing signals responsive to temperature near the catheter 110.

The control and delivery linkage 120 includes a metallic tube 223 defining a hollow lumen 224, and is further described with reference to figure 2.

5 In a preferred embodiment, the control and delivery linkage 120 is between about 1/2 inch (1.27 cm) and about 5/8 inches (1.59 cm) in diameter, and between about 6 inches (15.24 cm) and about 8 inches (20.32 cm) in length.

10 The control element 130 includes an electrode actuation element 131 for advancing the electrodes 113 out from the catheter 110, a electrode retraction element 132 for retracting the electrodes 113 into from the catheter 110, and an operation element 133 for controlling operation of the catheter 110, including delivery of flowable substances using the holes 112 and delivery of energy using the electrodes 113.

15 Advancing and Retracting Electrodes

Figure 2 shows a cut-away view of a catheter, taken along a line 2--2 in Figure 1.

20 The catheter 110 comprises a rotatable element 210 which is disposed for rotation in a first direction 211 to advance the electrodes 113 out of the catheter 110 and in a second direction 212 opposite the first direction 211 to retract the electrodes 113 back into the catheter 110.

25 In a preferred embodiment, the rotatable element 210 is coupled to a spring (not shown) or other device which holds the rotatable element 210 in a steady state with the electrodes 113 retracted into the catheter 110.

30 The rotatable element 210 is coupled to the electrode actuation element 131, which forces the rotatable element 210 to rotate in the first direction 211 so as to advance the electrodes 113 out of the catheter 110. When the actuator element is not actuated, the spring causes the rotatable element 210 to rotate in the second direction 212 so as to retract the electrodes 113 back into the catheter 110.

35 Each electrode 113 is coupled to an electrode carrier 220. In a preferred embodiment, each electrode carrier 220 is substantially bar-shaped (but is shown end-on in the figure) and is coupled to a plurality of electrodes 113, such

as about between about three and about six electrodes 113, so as to substantially simultaneously advance that plurality of electrodes 113 out of the catheter 110 and retract that plurality of electrodes 113 back into the catheter. A plurality of electrode carriers 220 are each disposed in a set of lines corresponding to lines of electrodes 113 disposed for advancement out of the catheter 110 and retraction back into the catheter 110.

In a preferred embodiment, the electrodes 113 may be disposed so that when advanced, the electrodes 113 extend to selected depths within the body structure to be ablated. These selected depths may be the same depth for all electrodes 113 which are advanced, or may include a first depth for a first set of electrodes 113 and a second depth for a second set of electrodes 113.

In a preferred embodiment, the electrode carriers 220 are coupled to a set of controls (not shown) in the control element 130 for selecting one or more electrode carriers 220 independently using one or more actuation levers 221, so as to be able to independently advance one or more sets of electrodes 113 coupled thereto out of the catheter 110 and to independently retract one or more sets of electrodes 113 back into the catheter 110.

Each electrode carrier 220 is coupled to the rotatable element 210 using a bearing 222, in such manner so as to translate rotation of the rotatable element 210 into linear radial movement of the electrodes 113. When the rotatable element 210 is rotated in the first direction 211, the electrodes are advanced in a first linear movement 223, while when the rotatable element 210 is rotated in the second direction 212, the electrodes are retracted in a second linear movement 224.

An interior 230 of the rotatable element 210 includes a lumen 225 through which fluids and other flowable substances are provided, and in which conductors providing control signals and sensor signals are disposed.

Operation of the Catheter and Electrode Assembly

Operation of the catheter and electrode assembly 100 includes at least the following steps:

- o The catheter 110 is inserted into the body at an opening, such as the rectum.

In a preferred embodiment, the opening is the rectum. A region of the rectum is first infused with a lubricant, such as K-Y jelly, and with an anesthetic, such as lidocaine. An anti-inflammatory, antispasmodic, or other condign medication would also be applied as appropriate. Thereafter, the catheter 110 is inserted into the lubricated region of the rectum. Due to the potential pain induced by the presence of the catheter 110 or electrodes 113, during operation the catheter 110 infuses a mixture of saline and lidocaine into the region 140 to be ablated.

In alternative embodiments, the opening may be another opening into the body, such as a natural orifice such as the vagina or the urethra, or an opening which has been made surgically, such as an incision which allows the catheter 110 to be inserted into a blood vessel.

The preferred size of the catheter 110 will of course be responsive to the size of the opening if other than the rectum. The choice of medicinal elements to be infused prior to or coeval with the catheter 110 will of course be responsive to judgments by medical personnel, and may include lubricants, anesthetics, antispasmodics, anti-inflammatories, antibiotics, or other materials with bioactive, chemoactive, or radioactive effect.

- o The catheter 110 is positioned within the body at a selected orientation and location, such as a position near a hemorrhoid.

In one preferred embodiment, the catheter 110 is positioned in the rectum near an external or internal hemorrhoid, in a manner as shown in figure 3. In this preferred embodiment, the electrodes 113 are ultimately advanced into the hemorrhoid to ablate the hemorrhoid.

In another preferred embodiment, the catheter 110 is positioned in the rectum near a prolapsed or spasmodic muscle, in a manner as shown in figure 4. In this preferred embodiment, the electrodes 113 are ultimately advanced into the prolapsed or spasmodic muscle to ablate selected portions of the prolapsed or spasmodic muscle.

5 In another preferred embodiment, the catheter 110 is positioned in the rectum near an anal fissure, in a manner as shown in figure 5. In this preferred embodiment, collagen is deposited into the fissure and the electrodes 113 are ultimately advanced into a region near the collagen to harden the collagen for filling the fissure.

10 In another preferred embodiment, the catheter 110 is positioned in the colon near a polyp, in a manner similar to that shown in figure 3. In this preferred embodiment, the electrodes 113 are ultimately advanced into the polyp to ablate the polyp.

15 In another preferred embodiment, the catheter 110 is positioned in the rectum near a pilonidal cyst, in a manner similar to that shown in figure 3. In this preferred embodiment, the electrodes 113 are ultimately advanced into the cyst to ablate the cyst.

20 In another preferred embodiment, the catheter 110 is positioned in the rectum, colon, large intestine, or small intestine, near a cyst or tumor, in a manner similar to that shown in figure 3. In this preferred embodiment, the electrodes 113 are ultimately advanced into the cyst or tumor to ablate the cyst or tumor.

25 In another preferred embodiment, the catheter 110 is positioned in a male patient, in the rectum near the prostate, in a manner as shown in figure 6. In this preferred embodiment, the electrodes 113 are ultimately advanced into a tumor in the prostate to ablate the tumor.

30 In another preferred embodiment, the catheter 110 is positioned in a female patient, in the vagina, near a cyst or fibroid, in a manner similar to that shown in Figure 3. In this preferred embodiment, the electrodes 113 are ultimately advanced into the cyst or fibroid to ablate the cyst or fibroid.

35 In another preferred embodiment, the catheter 110 is positioned in a female patient, in the vagina, near a prolapsed uterus, in a manner similar to that shown in Figure 4. In this preferred embodiment, the electrodes 113 are ultimately advanced into the prolapsed uterus selected portions of the prolapsed uterus.

- o The catheter 110 is anchored into place at the selected orientation and location by inflating a balloon 116, such as the distal balloon 116 and the proximal balloon 116.

5 In embodiments where the catheter 110 is positioned in the rectum, the catheter 110 is anchored into place using the proximal balloon 116 and the proximal balloon 116 operates in similar manner as a Foley catheter.

10 In alternative embodiments, the catheter 110 includes a stop balloon 116, such as a ring balloon (as shown in figure 3), disposed outside the body so as to prevent the catheter 110 from being inserted "too far", i.e., beyond its selected location.

- 15 o The region 140 near the catheter 110 is isolated from the rest of the body by inflating the distal balloon 116 and the proximal balloon 116. In a preferred embodiment, this step uses the same distal balloon 116 and the proximal balloon 116 as the step of anchoring the catheter 110 into place.

20 Isolation of the region 140 near the catheter 110 from the rest of the body need not be absolute. In a preferred embodiment, the distal balloon 116 and the proximal balloon 116 are microporous, are inflated using saline or water, and thus are disposed to provide saline or water into the region 140 near the catheter 110. However, in such an embodiment, gas and fluids from the rest of the body are allowed to leak into one or more of the balloons 116 and from there are allowed to
25 leak into the region 140 near the catheter 110.

30 Moreover, while in a preferred embodiment the seal made with the wall 141 of the region 140 by the balloon 116 is gas-tight, in alternative embodiments, that seal is allowed to be simply fluid-tight, and might allow gas to leak from the rest of the body into the region 140 near the catheter 110.

- 35 o One or more sets of electrodes 113 are selected for advancement into a selected mass of tissue in the region 140. The rotatable element 210 is rotated in the first direction 211, causing the selected sets of electrodes 113 to advance out of the catheter 110 and into the selected mass of tissue.

The selected set of electrodes 113 are just those electrodes 113 which are needed to penetrate the selected mass of tissue for ablation.

5 In a preferred embodiment where the selected mass of tissue for ablation is a hemorrhoid, the selected set of electrodes 113 are just those electrodes 113 which are needed to penetrate the hemorrhoid. If a plurality of hemorrhoids are selected for ablation, either (1) electrodes 113 needed to penetrate the plurality of hemorrhoids are selected, or (2) electrodes 113 needed to penetrate one of the hemorrhoids are selected, and the operation is repeated for each individual one of
10 the hemorrhoids.

Similarly, in preferred embodiments where the selected body structure for ablation is an individual cyst, fibroid, polyp, or tumor, the selected set of electrodes 113 are just those electrodes 113 which are needed to penetrate the
15 selected body structure. If there is more than one such selected body structure, either (1) more than one set of electrodes 113 may be selected, or (2) just one set of electrodes 113 may be selected and the operation is repeated for each individual such body structure.

20 Similarly, in preferred embodiments where the selected body structure for ablation is muscle tissue or other tissue which is part of a larger body structure, such as a prolapsed or spasmodic muscle, the selected set of electrodes 113 are just those one or more sets of electrodes 113 which are needed to penetrate the portion of the body structure which has been selected for ablation.

25 o Flowable substances are provided using the holes 112, and energy is provided to the electrodes 113, so as to ablate the mass of tissue in the region 140.

30 In a preferred embodiment, the flowable substances are provided using the holes 112 to the region 140 near the catheter 110.

35 In alternative embodiments, the flowable substances may be provided, in addition or instead, (1) from an area of the catheter covered by a microporous membrane, or (2) from one or more microporous balloons. The microporous balloons may either be the same as or in addition to the balloons 116 used to anchor the catheter in place or to block gas or fluid.

In preferred embodiments, the flowable substances have one of the following functions: (1) to aid in ablation, such as by transmitting RF energy from the electrodes 113 to the body structure to be ablated, as is done by saline or other electrolytic solutions, (2) to rehydrate tissue, as in done by saline or water, or (3) to repair tissue, such as by flowing into cysts or fissures or voids, or by covering lesions, as is done by collagen in a soft form which can be hardened by RF energy.

In a preferred embodiment, the electrodes 113 deliver RF energy having a frequency between about 435 megahertz and about 485 megahertz, for a period between about 5 minutes and about 10 minutes. The RF energy is received by and heats tissue and other body structures near the electrodes 113, causing ablation by means of cell death, dehydration, or denaturation.

In alternative embodiments, the electrodes 113 may deliver other forms of energy, such as heat, microwaves, or infrared or visible laser energy.

The electrodes 113 are controlled by a feedback technique, using the at least one sensor 119. In embodiments where there is more than one sensor 119, the feedback technique may be responsive to each sensor 119.

In one preferred embodiment, the at least one sensor 119 includes a temperature sensor 119 and the feedback technique includes a microprocessor (not shown) disposed in or coupled to the control element 130 and operating under control of application software for maintaining the temperature of the body structure to be ablated at a selected temperature, such as a temperature exceeding between about 90° Celsius and about 120° Celsius. In this preferred embodiment, the microprocessor also controls delivery of fluids for cooling or hydration, so as to maintain the temperature of surrounding tissue (i.e., other than the tissue selected for ablation) at temperatures less than between about 90° Celsius and about 120° Celsius.

In another preferred embodiment, the at least one sensor 119 also includes an impedance sensor 119 and the feedback technique includes a microprocessor operating to terminate delivery of RF energy when a measured

impedance of the body structure to be ablated undergoes a substantial change indicative of dehydration or denaturation.

- 5 o One or more sets of electrodes 113 are selected for retraction back from the selected mass of tissue in the region 140. The rotatable element 210 is rotated in the second direction 211, causing the selected sets of electrodes 113 to retract out of the selected mass of tissue and back into the catheter 110.

10 The same electrodes 113 which were advanced out of the catheter 110 are retracted back into the catheter 110.

- o The catheter 110 is withdrawn from the body at the opening through which it was inserted.

15 Before removal, the balloons 116 are deflated so the catheter 110 is no longer anchored in place, all electrodes 113 are retracted back into the catheter 110, and the catheter 113 is configured to no longer provide flowable substances or energy for ablation.

20 Particular Methods and Apparatus for Treatments

In preferred embodiments, the catheter and electrode assembly 100 may also be used for treatments in addition to, or instead of, ablation of body structures or tissue.

25

In one preferred embodiment, operation of the catheter and electrode assembly 100 includes at least the following steps:

- 30 o The catheter 110 is inserted into a natural body lumen, such as the urethra.

In a preferred embodiment, the natural body lumen comprises a normally tubular body structure which has prolapsed, is spasmodic, or is otherwise subject to blockage (partial or complete) or damage (such as to a wall of the natural body lumen).

35

- o The catheter 110 infuses a hardenable substance into the natural body lumen, so as to coat at least one selected section of the wall of the natural body lumen.

5 In a preferred embodiment, the hardenable substance includes a collagen which is capable of being flowed onto the wall of the natural body lumen and which is capable of being hardened by application of RF energy, heat, or another agent to be provided by the catheter and electrode assembly 100.

- 10 o The electrodes 113 are advanced and deliver energy to the hardenable substance to harden it.

In a preferred embodiment, the holes 112 provide saline and the electrodes 113 deliver RF energy to the collagen to harden it, so as to form a hard covering to the wall of the natural body lumen. If appropriate, more than one layer of collagen is applied, so as to provide a hard covering having a thickness exceeding a selected threshold, such as 0.1 inch (0.25 cm). The particular selected threshold will of course depend on the preferred diameter of the natural body lumen.

20 In a preferred embodiment for treatment of a prolapsed or spasmodic muscle, (1) the catheter 110 is inserted and pushed through a region where the muscle has prolapsed or blocked the rectum, colon, large intestine, or small intestine, (2) the prolapsed or spasmodic muscle is partially ablated, and (3) collagen is infused and hardened to strengthen the muscle wall. In alternative embodiments, the collagen may be infused before ablation in one or more boluses deposited within the muscle (or on the muscle or near the muscle), so that the steps of muscle ablation and collagen hardening will occur substantially simultaneously.

30 In a preferred embodiment for treatment of an anal fissure, (1) the catheter 110 is inserted into a region where the fissure has occurred, (2) a suspension of collagen and saline is infused and fills the fissure, and (3) the collagen is hardened while the saline is removed from the suspension. In this preferred embodiment, the isolated region between the distal balloon 116 and the proximal balloon 116 is maintained at a positive differential pressure with respect to the rest of the rectum, so that the collagen infuses into the fissure; this procedure or a similar procedure is also followed for treatment of diverticulosis and diverticulitis.

In a preferred embodiment for treatment, in a female patient, of a prolapsed uterus, (1) the catheter 110 is inserted into a region where the uterus has prolapsed, (2) the prolapsed uterus is partially ablated, and (3) collagen is infused and hardened to strengthen the muscle wall. Similarly to treatment of a prolapsed muscle, in alternative embodiments, the collagen may be infused before ablation, so that the steps of muscle ablation and collagen hardening will occur substantially simultaneously.

10 Alternative Embodiments

Although preferred embodiments are disclosed herein, many variations are possible which remain within the concept, scope, and spirit of the invention, and these variations would become clear to those skilled in the art after perusal of this application.

15

Claims

We claim:

1. A method for ablating a structure within a body, said method including the steps of
5 inserting a catheter into a region of said body;
isolating said region from gas or fluid from outside said region;
emitting, from an electrode coupled to said catheter into said region, an effective amount of energy to ablate said structure.
- 10 2. A method as in claim 1, wherein said region is in or near the sphincter, rectum, colon, or prostate.
3. A method as in claim 1, wherein said structure includes a cyst, hemorrhoid, polyp, tumor, or lesion.
15
4. A method as in claim 1, wherein said energy includes RF energy having a frequency between about 435 megahertz and about 485 megahertz.
5. A method as in claim 1, wherein said step of emitting is
20 conducted for less than about 10 minutes.
6. A method as in claim 1, including the step of advancing said electrode from within said catheter to a point inside a selected mass of tissue within said region.
25
7. A method as in claim 1, wherein said step of isolating includes the step of blocking gas or fluid from entering said region using at least one balloon.
8. A method as in claim 7, wherein said at least one balloon
30 includes a balloon disposed at an upstream location or disposed at a downstream locations from said region.
9. A method as in claim 7, wherein said step of isolating includes the step of inflating said at least one balloon.
35

10. A method as in claim 7, including the step of anchoring said catheter in a selected location using said at least one balloon.
11. A method as in claim 1, including the step of anchoring said catheter in a selected location.
12. A method as in claim 11, wherein said step of anchoring includes the step of disposing at least one balloon against a wall of said region.
13. A method as in claim 1, including the steps of sensing at least one control parameter of said region; and controlling said step of emitting using a feedback technique including said control parameter.
14. A method as in claim 1, including the step of delivering a flowable substance to said region.
15. A method as in claim 14, including the steps of sensing at least one control parameter of said region; and controlling said step of delivering using a feedback technique including said control parameter.
16. A method as in claim 14, wherein said flowable substance includes an effective amount of fluid to cool said region to a selected temperature.
17. A method as in claim 14, wherein said flowable substance includes an effective amount of fluid to rehydrate a selected mass of tissue in said region.
18. A method as in claim 14, wherein said flowable substance includes saline.
19. A method as in claim 14, wherein said flowable substance includes an antibiotic.

20. A method as in claim 14, wherein said flowable substance includes collagen.

21. A method as in claim 14, wherein said flowable substance
5 includes a substance for covering or filling a selected structure within said region.

22. A method as in claim 14, wherein said step of delivering includes exuding said flowable substance from at least one hole in said catheter.

10 23. A method as in claim 14, wherein said step of delivering includes exuding said flowable substance from a microporous membrane.

24. A method as in claim 14, wherein said step of delivering includes exuding said flowable substance from a balloon having a microporous membrane.
15

25. A method as in claim 1, including the step of coupling light from said region to a point outside the body, whereby an operator is able to view said region.

20 26. A method as in claim 1, including the steps of
receiving light from said region representative of a view of said region;
and
controlling an orientation or position of said catheter responsive to said
view.
25

27. Apparatus for ablating a structure in a region within a body, said
apparatus including
a catheter disposed for positioning in said region;
means for isolating said region from gas or fluid from outside said
30 region; and
an electrode coupled to said catheter.

28. Apparatus as in claim 27, wherein said region is in or near the
sphincter, rectum, colon, or prostate.
35

29. Apparatus as in claim 27, wherein said structure includes a cyst, hemorrhoid, polyp, tumor, or lesion.

30. Apparatus as in claim 27, wherein said energy includes RF
5 energy having a frequency between about 435 megahertz and about 485 megahertz.

31. Apparatus as in claim 27, including means for advancing said electrode from within said catheter to a point inside a selected mass of tissue within said region.

10

32. Apparatus as in claim 27, wherein said means for isolating includes at least one balloon.

33. Apparatus as in claim 32, wherein said at least one balloon
15 includes a balloon disposed at an upstream location or disposed at a downstream locations from said region.

34. Apparatus as in claim 32, including means for inflating said at least one balloon.

20

35. Apparatus as in claim 32, wherein said at least one balloon is operative to anchor said catheter in a selected location.

36. Apparatus as in claim 27, including means for anchoring said
25 catheter in a selected location.

37. Apparatus as in claim 36, wherein said means for anchoring includes at least one balloon capable of being disposed against a wall of said region.

30

38. Apparatus as in claim 27, including at least one control parameter sensor; and means, responsive to said control parameter sensor, for controlling said electrode.

35

39. Apparatus as in claim 27, including means for delivering a flowable substance to said region.

40. Apparatus as in claim 39, including
at least one control parameter sensor; and
means, responsive to said control parameter sensor, for controlling said
5 means for delivering.

41. Apparatus as in claim 39, wherein said flowable substance
includes an effective amount of fluid to cool said region to a selected temperature.

10 42. Apparatus as in claim 39, wherein said flowable substance
includes an effective amount of fluid to rehydrate a selected mass of tissue in said
region.

43. Apparatus as in claim 39, wherein said flowable substance
15 includes saline.

44. Apparatus as in claim 39, wherein said flowable substance
includes an antibiotic.

20 45. Apparatus as in claim 39, wherein said flowable substance
includes collagen.

46. Apparatus as in claim 39, wherein said flowable substance
includes a substance for covering or filling a selected structure within said region.
25

47. Apparatus as in claim 39, wherein said means for delivering
includes at least one hole in said catheter.

48. Apparatus as in claim 39, wherein said means for delivering
30 includes a microporous membrane.

49. Apparatus as in claim 39, wherein said means for delivering
includes a balloon having a microporous membrane.

35 50. Apparatus as in claim 27, including an optical path from said
region to a point outside the body, whereby an operator is able to view said region.

51. Apparatus as in claim 50, including means for controlling an orientation or position of said catheter responsive to said view.
- 5 52. Apparatus as in claim 50, including a view piece coupled to said optical path.

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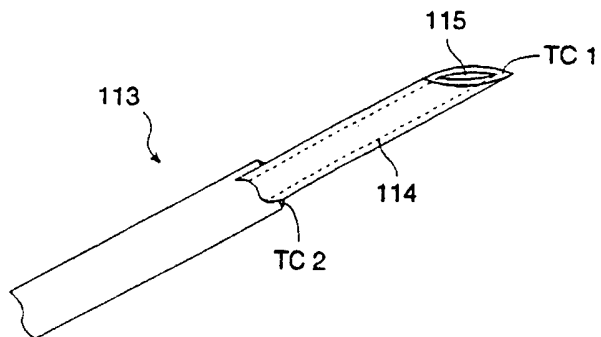


FIG. 1A

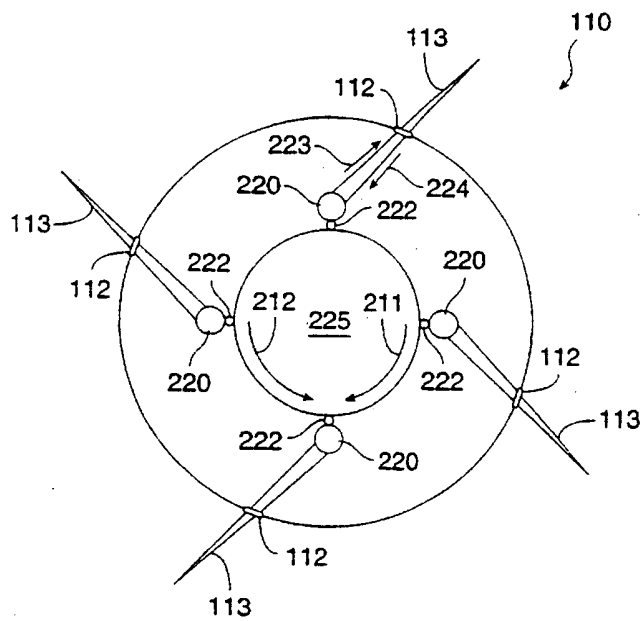


FIG. 2

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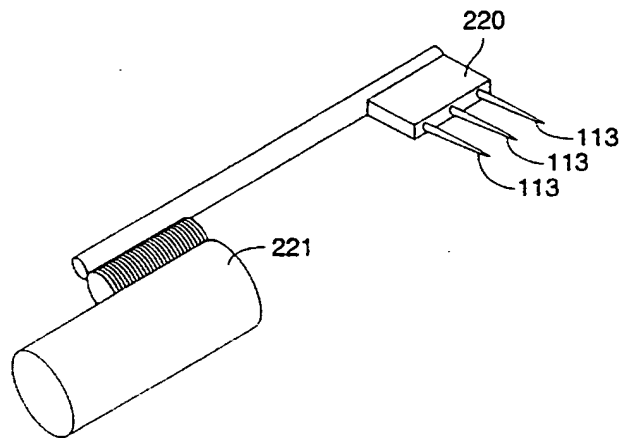


FIG. 2A

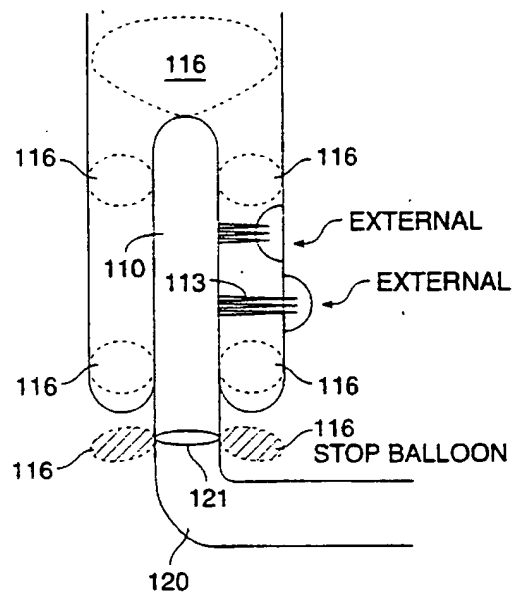


FIG. 3

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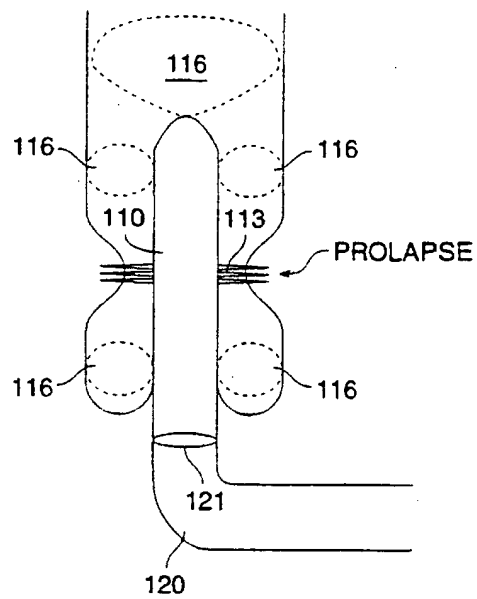


FIG. 4

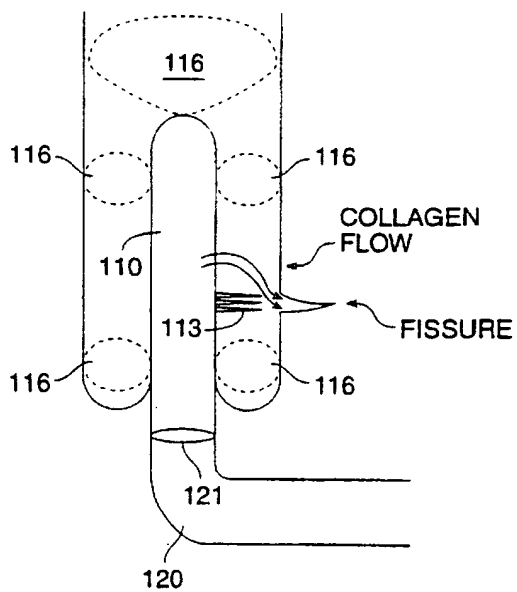


FIG. 5

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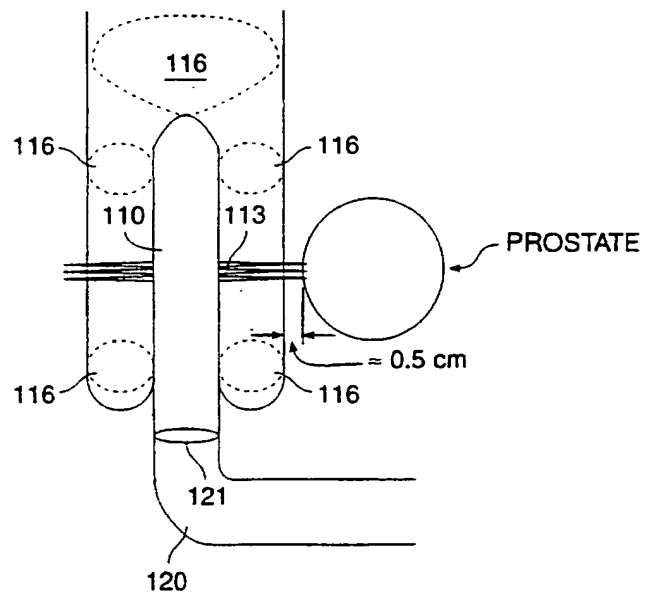


FIG. 6

SUBSTITUTE SHEET (RULE 26)

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 97/16506

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 A61B17/39

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 A61B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GB 2 269 538 A (VIDAMED) 16 February 1994 see the whole document ---	27-52
A	US 5 342 357 A (NARDELLA) 30 August 1994 see abstract; figure 12 ---	39-44
A	US 5 458 568 A (RACCHINI) 17 October 1995 see abstract; figures 3,5 ---	49
P,A	US 5 569 241 A (EDWARDS) 29 October 1996 see column 6, line 35 - line 40 -----	47-52

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
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Date of the actual completion of the international search

11 February 1998

Date of mailing of the international search report

23.02.98

Name and mailing address of the ISA

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INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 97/16506

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons.

1. ☒ Claims Nos.: 1-26
because they relate to subject matter not required to be searched by this Authority, namely:
Rule 39.1(iv) PCT - Method for treatment of the human or animal body by surgery
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 97/16506

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Information on patent family members

International Application No

PCT/US 97/16506

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Information on patient family members

PCT/US 97/16506

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